

BIOGRAPHICAL SKETCH

NAME		POSITION TITLE		
Sandro Banfi, M.D.		Researcher, TIGEM		
INSTITUTION AND LOCATION		DEGREE	YEAR CONFERRED	FIELD OF STUDY
Federico II University, Naples, Italy		MD	1989	Medicine
Federico II University, Naples, Italy		Residency	1993	Neurology

Research and Professional Experience

1988-1989	Predoctoral Fellow in Neurology at the Institute of Neurology, School of Medicine, Federico II University, Naples, Italy; Director: Prof. G. Campanella; Supervisor: Dr. A. Filla.
1989-1991	Postdoctoral Fellow in Neurology at the Institute of Neurology, School of Medicine, Federico II University, Naples, Italy; Director: Prof. G. Campanella; Supervisor: Dr. A. Filla.
03/91-08/91	Postdoctoral Fellow at the Institute for Molecular Genetics, School of Medicine, Federico II University, Naples, Italy; Supervisor: Dr. S. Cocozza.
09/91-04/92	Postdoctoral Fellow at the Department of Pediatrics, Baylor College of Medicine, Houston, TX; Professor and Director: R.D. Feigin; Supervisor: Dr. H.Y. Zoghbi.
05/92-12/94	Research Associate at the Department of Pediatrics, Baylor College of Medicine, Houston, TX; Professor and Director: R.D. Feigin; Supervisor: Dr. H.Y. Zoghbi.
1995-Present	Researcher , Telethon Institute of Genetics and Medicine (TIGEM), Milan, Italy; Director: Prof. A. Ballabio.
10/04-present	Lecturer , European School of Molecular Medicine (SEMM), Milan and Naples, Italy.

Publications

Filla A, De Michele G, Marconi R, Santorelli F, Trombetta L, **Banfi S**, Campanella G: Effects of thyrotropin-releasing hormone on heart rate in inherited ataxias. Med Sci Res 17: 569-570, 1989.

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Ranum LPW, Chung M-Y, **Banfi S**, Bryer A, Schut LJ, Ramesar R, Duvick LA, McCall AE, Subramony SH, Goldfarb L, Gomez C, Sandkuijl LA, Orr HT, Zoghbi HY: Molecular and clinical correlations in spinocerebellar ataxia type 1 (SCA1): Evidence for

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Banfi S, Servadio A, Chung M-y, Capozzoli F, Duvick L, Elde R, Zoghbi HY, Orr HT: Cloning and developmental expression analysis of the murine homolog of the spinocerebellar ataxia type 1 gene (Sca1). Human Molecular Genetics 5:33, 1996.

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Book chapters

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